

WHAT IS CLAIMED IS:

1	 A substantially purified immunoglobulin
2	polypeptide that specifically binds to a human type beta
3	platelet-derived growth factor receptor (βPDGF-R), wherein
4	binding of the polypeptide has one or more of the following
5	effects:
6	i) inhibition of PDGF BB or AB binding to the
7	βPDGF-R;
8	ii) inhibition of PDGF-induced β PDGF-R
9	phosphorylation;
10	iii) inhibition of PDGF-induced dimerization of
11	β PDGF-R;
12	iv) inhibition of PDGF-induced mitogenesis of
13	cells displaying human β PDGF-R; and
14	v) inhibition of PDGF-induced chemotaxis and
15	migration of cells displaying β PDGF-R.

- 2. An immunoglobulin polypeptide of claim 1, wherein the polypeptide is a monoclonal antibody.
- 3. An immunoglobulin polypeptide of claim 2, wherein the monoclonal antibody is antibody 2A1E2.
- 4. A substantially purified polypeptide having an amino acid sequence substantially identical to a sequence of a complementarity determining region of an immunoglobulin polypeptide of claim 1.
- 5. An immunoglobulin polypeptide of claim 1, wherein the polypeptide is linked to a detectable label.
- 6. An immunoglobulin polypeptide of claim 1, wherein the polypeptide is chimeric.
- 7. A substantially purified immunoglobulin polypeptide that specifically recognizes an epitope which lies

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- in the second Ig-like domain in the extracellular region of the β PDGF-R.
- 8. A composition comprising a monoclonal antibody or binding fragment thereof that binds to the human β PDGF-R, which antibody or fragment inhibits <u>in vivo</u> binding of PDGF BB or AB to the receptor.
- 9. An isolated nucleic acid having a sequence
 substantially identical to a nucleic acid coding for an
 immunoglobulin polypeptide or a binding fragment thereof,
 wherein binding of the polypeptide or fragment to a human
 βPDGF-R has one or more of the following effects:
 - i) inhibition of PDGF BB or AB binding to the β PDGF-R;
- 8 ii) inhibition of PDGF-induced β PDGF-R phosphorylation;
- iii) inhibition of PDGF-induced dimerization of
 βPDGF-R;
- iv) inhibition of PDGF-induced mitogenesis of cells displaying the human β PDGF-R; and
- 14 v) inhibition of PDGF-induced chemotaxis and 15 migration of cells displaying β PDGR-R.
- 1 10. A nucleic acid of claim 9, wherein the nucleic 2 acid is operably linked to a promoter.
- 1 11. A nucleic acid of claim 10, wherein the promoter 2 and the nucleic acid are contained in an expression vector.
- 1 12. A cell line transfected, transformed, or infected with a nucleic acid of claim 9.
- 13. A method of producing a substantially purified 2 immunoglobulin polypeptide, or binding fragment thereof, which 3 binds to a human type beta PDGF receptor (β PDGF-R), wherein the 4 binding of the polypeptide or fragment to the β PDGF-R has one

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- or more of the following effects: inhibition of PDGF BB or AB 5
- binding to the β PDGF- \hat{R} ; inhibition of PDGF-induced β PDGF-R 6
- phosphorylation; inhibition of PDGF-induced dimerization of the 7 .
- 8 βPDGF-R; and inhibition of PDGF-induced mitogenesis of cells
- displaying human β PDGF-R; and inhibition of PDGF-induced 9
- 10 chemotaxis and migration\of cells displaying human β -PDGF; the
- 11 method comprising:
- 12 i) growing a cell line comprising a nucleic acid
- 13 encoding the immunoglobulin polypeptide; and
- harvesting the immunoglobulin polypeptide. 14 ii)
 - A method of claim 13, wherein the cell line is a 1 14. 2 hybridoma.
- 15. A method of claim 14, wherein the hybridoma is 2 ATCC no. HB10938.
- A method of claim 13, wherein the immunoglobulin 1 16. polypeptide is a monoclonal antibody. 2
- A method of treating a human having a 17. PDGF-mediated disease involving proliferation, migration or chemotaxis of smooth muscle cells, comprising administering to the patient a therapeutically effective dose of at least one 54 immunoglobulin polypeptide according to claim 1, or fragments of the immunoglobulin polypeptide, and a pharmaceutically acceptable carrier.
- An isolated cell line designated as ATCC no. 1
- 2 HB10938.

and and